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**INFORMATION DISCLOSURE  
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Application Number	10/806,295
Filing Date	March 22, 2004
First Named Inventor	LaColla, <i>et al.</i>
Group Art Unit	Unassigned
Examiner Name	Unassigned
Attorney Docket Number	06171.105033 IDX 1008 DIV

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**U.S. PATENT DOCUMENTS**

Examiner Initials *	Cite No. <sup>1</sup>	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number	Kind Code (if known)			
Gm	AA	4,866,084	A	Gunasekera <i>et al.</i>	09-12-1989	
	AB	5,124,327	A	Greenlee <i>et al.</i>	06-23-1992	
	AC	5,489,685	A	Houpis <i>et al.</i>	02-06-1996	
	AD	5,527,819	A	Williams <i>et al.</i>	06-18-1996	
	AE	5,830,894	A	Pevear <i>et al.</i>	11-03-1998	
	AF	5,852,011	A	Matsunaga <i>et al.</i>	12-22-1998	
	AG	5,929,114	A	Domagala <i>et al.</i>	07-27-1999	
	AH	5,935,982	A	Dykstra <i>et al.</i>	08-10-1999	
	AI	5,945,440	A	Kleinschroth <i>et al.</i>	08-31-1999	
	AJ	5,981,525	A	Farina <i>et al.</i>	11-09-1999	
	AK	6,025,390	A	Farina <i>et al.</i>	02-15-2000	

**FOREIGN PATENT DOCUMENTS**

Examiner Initials *	Cite No. <sup>1</sup>	Foreign Patent Document			Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T <sup>6</sup>
		Office <sup>3</sup>	Number	Kind Code <sup>2</sup> (if known)				
Gm	AL	EP	0,530,907	A1	Merck & Co.	03-10-1993		
	AM	WO	94/19321	A1	Merck & Co., Theoharides	09-01-1994		
Gm	AN	WO	02/083126	A1	Idenix; Univ. Degli Studi di Cagliari	10-24-2002		
Gm	AO	WO	04/014364	A1	Idenix	02-19-2004		

**OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS**

Examiner Initials *	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>6</sup>
Gm	AP	BALANI, S.K., <i>et al.</i> , "Biotransformation of 5-chloro-3-phenylthioindole-2-carboxamide (L-734,005) in rhesus monkeys and rat liver microsomes to a potent HIV-1 reverse transcriptase inhibitor," <i>Drug Metab. Dispos.</i> , 21(4):598-604 (July-August 1993).	

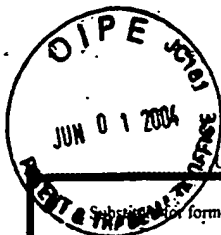
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	<del>AQ</del>	<del>CLAUSON-KAAS, N., et al., "Preparation of Cis and Trans 2,5-Dimethoxy-2-(acetamidomethyl)-2,5-dihydrofuran, of Cis and Trans 2,5-Dimethoxy-2-(acetamidomethyl)-tetrahydrofuran and of 1-Phenyl-2-(acetamidomethyl)pyrrole," Acta Chem. Scand., 6:667-670 (1952).</del>	
	<del>BA</del>	<del>ELMING, N., et al., "The preparation of pyrroles from furans," Acta Chem. Scand., 6:867-874 (1952).</del>	
	<del>BB</del>	<del>GAGLIARDI, S., et al., "5-(5,6-Dichloro-2-indolyl)-2-methoxy-2,4-pentadienamides: novel and selective inhibitors of the vacuolar H<sup>+</sup>-ATPase of osteoclasts with bone antiresorptive activity," J. Med. Chem., 41(10):1568-1573 (May 7, 1998).</del>	
	<del>BC</del>	<del>PAUWELS, R., et al., "Potent and selective inhibition of HIV-1 replication in vitro by a novel series of TIBO derivatives," Nature, 343(6257):470-474 (February 1, 1990).</del>	
	<del>BD</del>	<del>PAUWELS, R., et al., "Potent and highly selective human immunodeficiency virus type 1 (HIV-1) inhibition by a series of alpha-anilinophenylacetamide derivatives targeted at HIV-1 reverse transcriptase," Proc. Natl. Acad. Sci. USA, 90(5):1711-1715 (March 1, 1993).</del>	
	<del>BE</del>	<del>PHILLIPS, R.R., "The Japp-Klingemann Reaction," Org. Reactions, 10:143-178 (1959).</del>	
	<del>BF</del>	<del>ROMERO, D.L., et al., "Bis(heteroaryl)piperazine (BHAP) reverse transcriptase inhibitors: structure-activity relationships of novel substituted indole analogues and the identification of 1-[(5-methanesulfonamido-1H-indol-2-yl)-carbonyl]-4-[3-[(1-methylethyl)amino]-pyridinyl]piperazine monomethanesulfonate (U-90152S), a second-generation clinical candidate," J. Med. Chem., 36(10):1505-1508 (May 14, 1993).</del>	
	<del>BG</del>	<del>WILLIAMS, T.M., et al., "5-chloro-3-(phenylsulfonyl)indole-2-carboxamide: a novel, non-nucleoside inhibitor of HIV-1 reverse transcriptase, J. Med. Chem., 36(9):1291-1294 (April 30, 1993).</del>	

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